Belinda Gray

List of Publications by Year
in descending order

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33 papers	744 citations	567281 15 h-index	526287 27 g-index
34 all docs	34 docs citations	34 times ranked	1288 citing authors

#	Article	IF	Citations
1	Rare Variation in Drug Metabolism and Long QT Genes and the Genetic Susceptibility to Acquired Long QT Syndrome. Circulation Genomic and Precision Medicine, 2022, 15, CIRCGEN121003391.	3.6	7
2	â€œConcealed cardiomyopathyâ€•as a cause of previously unexplained sudden cardiac arrest. International Journal of Cardiology, 2021, 324, 96-101.	1.7	37
3	Comparison of conventional autopsy with post-mortem magnetic resonance, computed tomography in determining the cause of unexplained death. Forensic Science, Medicine, and Pathology, 2021, 17, 10-18.	1.4	13
4	Biventricular Myocardial Fibrosis and Sudden Death in Patients With BrugadaÂSyndrome. Journal of the American College of Cardiology, 2021, 78, 1511-1521.	2.8	18
5	Accuracy of the 2017 international recommendations for clinicians who interpret adolescent athletesâ€™ ECGs: a cohort study of 11 168 British white and black soccer players. British Journal of Sports Medicine, 2020, 54, 739-745.	6.7	41
6	Genetic Testing for Inherited Cardiovascular Disease: Implications of the AHA Scientific Statement for Cardiologists. Heart Lung and Circulation, 2020, 29, 1581-1584.	0.4	1
7	Prevalence and Phenotypic Correlations of Calmodulinopathy-Causative CALM1-3 Variants Detected in a Multicenter Molecular Autopsy Cohort of Sudden Unexplained Death Victims. Circulation Genomic and Precision Medicine, 2020, 13, e003032.	3.6	3
8	When do athletes benefit from cardiac genetic testing?. British Journal of Sports Medicine, 2020, 54, 939-940.	6.7	1
9	Utility of genetic testing in athletes. Clinical Cardiology, 2020, 43, 915-920.	1.8	11
10	Triadin Knockout Syndrome Is Absent in a Multi-Center Molecular Autopsy Cohort of Sudden Infant Death Syndrome and Sudden Unexplained Death in the Young and Is Extremely Rare in the General Population. Circulation Genomic and Precision Medicine, 2020, 13, e002731.	3.6	4
11	Patients With Genetic Heart Disease and COVID-19: A Cardiac Society of Australia and New Zealand (CSANZ) Consensus Statement. Heart Lung and Circulation, 2020, 29, e85-e87.	0.4	4
12	Electrocardiography in Athletes â€“ How to Identify High-risk Subjects. European Journal of Arrhythmia & Electrophysiology, 2020, 6, 24.	0.2	1
13	Evaluation After Sudden Death in the Young. Circulation: Arrhythmia and Electrophysiology, 2019, 12, e007453.	4.8	19
14	Sudden Death and Left Ventricular Involvement in Arrhythmogenic Cardiomyopathy. Circulation, 2019, 139, 1786-1797.	1.6	122
15	A balanced translocation disrupting SCN5A in a family with Brugada syndrome and sudden cardiac death. Heart Rhythm, 2019, 16, 231-238.	0.7	13
16	Editorial commentary: Will the real long QT genes please stand up. Trends in Cardiovascular Medicine, 2018, 28, 465-466.	4.9	0
17	The Diagnostic Yield of Brugada Syndrome After Sudden Death WithÂNormal Autopsy. Journal of the American College of Cardiology, 2018, 71, 1204-1214.	2.8	84
18	Exome-Wide Rare Variant Analyses in Sudden Infant Death Syndrome. Journal of Pediatrics, 2018, 203, 423-428.e11.	1.8	17

#	Article	IF	Citations
19	Lack of genotype-phenotype correlation in Brugada Syndrome and Sudden Arrhythmic Death Syndrome families with reported pathogenic SCN1B variants. Heart Rhythm, 2018, 15, 1051-1057.	0.7	15
20	Relations between right ventricular morphology and clinical, electrical and genetic parameters in Brugada Syndrome. PLoS ONE, 2018, 13, e0195594.	2.5	23
21	Cardiovascular Effects of Energy Drinks in Familial Long QT Syndrome: A Randomized Cross-Over Study. International Journal of Cardiology, 2017, 231, 150-154.	1.7	35
22	Twelve-lead ambulatory electrocardiographic monitoring in Brugada syndrome: Potential diagnostic and prognostic implications. Heart Rhythm, 2017, 14, 866-874.	0.7	47
23	New Insights Into the Genetic Basis of Inherited Arrhythmia Syndromes. Circulation: Cardiovascular Genetics, 2016, 9, 569-577.	5.1	45
24	A novel heterozygous mutation in cardiac calsequestrin causes autosomal dominant catecholaminergic polymorphic ventricular tachycardia. Heart Rhythm, 2016, 13, 1652-1660.	0.7	60
25	Clinical and genetic features of Australian families with long QT syndrome: A registryâ€based study. Journal of Arrhythmia, 2016, 32, 456-461.	1.2	9
26	NOS1AP Polymorphisms Modify QTc Interval Duration But Not Cardiac Arrest Risk in Hypertrophic Cardiomyopathy. Journal of Cardiovascular Electrophysiology, 2015, 26, 1346-1351.	1.7	4
27	Radiation Exposure During Cardiac Catheterisation is Similar for Both Femoral and Radial Approaches. Heart Lung and Circulation, 2015, 24, 264-269.	0.4	13
28	Social determinants of health in the setting of hypertrophic cardiomyopathy. International Journal of Cardiology, 2015, 184, 743-749.	1.7	25
29	Brugada Syndrome: A Heterogeneous Disease with a Common ECG Phenotype?. Journal of Cardiovascular Electrophysiology, 2014, 25, 450-456.	1.7	18
30	Late positive flecainide challenge test for Brugada syndrome. Heart Rhythm, 2014, 11, 898-900.	0.7	5
31	Severe hypertensive encephalopathy following percutaneous balloon aortic valvuloplasty for aortic stenosis. International Journal of Cardiology, 2014, 171, e63-e64.	1.7	1
32	Prolongation of the QTc Interval Predicts Appropriate Implantable Cardioverter-Defibrillator Therapies in Hypertrophic Cardiomyopathy. JACC: Heart Failure, 2013, 1, 149-155.	4.1	37
33	Homozygous mutation in the cardiac troponin I gene: Clinical heterogeneity in hypertrophic cardiomyopathy. International Journal of Cardiology, 2013, 168, 1530-1531.	1.7	11